

## Amendments to the Claims

### **1-88. (Cancelled)**

**89. (Currently Amended)** A fast-dissolving pharmaceutical composition in a solid dosage form ~~excepting a nanoparticle suspension~~, comprising micronized (R)-2-(4-bromo-2-fluorobenzyl)-1,2,3,4-tetrahydropyrrolo[1,2-a]pyrazine-4-spiro-3'-pyrrolidine-1,2',3,5'-tetrone (hereinafter referred to as "AS-3201") having a mean particle size of in a range of ~~above 1~~ about 1.5  $\mu\text{m}$  to less than about 10  $\mu\text{m}$  in a ratio of about 0.5% by weight to about 25% by weight of the total weight of the pharmaceutical composition, and as a stabilizer at least one acidic substance having a pKa less than about 5.6,

wherein when a dissolution percentage of AS-3201 from the composition is measured according to the Paddle method, 50% or more of the AS-3201 in the composition is dissolved with 15 minutes from the start of the method.

**90. (Currently Amended)** The fast-dissolving pharmaceutical composition according to claim 89, wherein the acidic substance is ~~tartaric acid, a member selected from the group consisting of citric acid, tartaric acid, maleic acid and phosphoric acetate.~~

### **91-113. (Cancelled)**

**114. (Previously presented)** The fast-dissolving pharmaceutical composition according to claim 89, wherein the mean particle size of the micronized AS-3201 is in the range of about 1.5  $\mu\text{m}$  to about 5  $\mu\text{m}$ .

### **115. (Cancelled)**

**116. (Previously presented)** The fast-dissolving pharmaceutical composition according to claim 89, wherein the mean particle size of the micronized AS-3201 is in the range of about 1.5  $\mu\text{m}$  to about 3  $\mu\text{m}$ .

**117. (Previously presented)** The fast-dissolving pharmaceutical composition according to claim 89, wherein the solid dosage form is tablets, capsules, granules or powder.

**118. (New)** The fast-dissolving pharmaceutical composition according to claim 89, wherein the acidic substance is tartaric acid.